

Drug Enforcement Administration

Importer of Controlled Substances Application: Groff NA Hemplex, LLC

[Docket No. DEA-903]

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Groff NA Hemplex, LLC. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration,
Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield,
Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement
Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.
All request for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn:
Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug
Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701
Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on August 13, 2021, Groff NA Hemplex, LLC., 100 Redco Avenue, Suite A, Red Lion, Pennsylvania 17356-1436, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled Substance	Drug Code	Schedule
Marihuana Extract	7350	I
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Tetrahydrocannabinols	7370	I

The company plans to import the above controlled substances as bulk to manufacture research grade material for clinical trial studies. Several types of Marihuana Extract compounds are listed under drug code 7350. No other activity for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration- approved or non-approved finished dosage forms for commercial sale.

Brian S. Besser,

Acting Assistant Administrator.

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